Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (Original) A method for reducing the severity of a bone fracture in a subject, the method comprising administering to a site of said bone fracture in said subject a therapeutically effective amount of an agent that inhibits activity or expression of a BMP-3 polypeptide.
 - 2. (Original) The method of claim 1, wherein said agent is an anti-BMP-3 antibody.
- 3. (Original) The method of claim 2, wherein said antibody is a monoclonal antibody.
- 4. (Original) The method of claim 3, wherein said monoclonal antibody is a human monoclonal antibody or a humanized monoclonal antibody.
- 5. (Original) The method of claim 1, wherein said agent is an anti-BMP-3 antisense RNA.
 - 6. (Original) The method of claim 1, wherein said subject is a human.
- 7. (Original) The method of claim 1, wherein said agent is administered systemically to said subject.
 - 8. (Original) The method of claim 7, wherein said administration is intravenous.
- 9. (Original) The method of claim 1, wherein said agent is administered locally to said site.
- 10. (Original) The method of claim 9, wherein said agent is administered by intraosseous injection.

- 11. (Original) The method of claim 1, wherein said agent is administered in conjunction with a matrix.
- 12. (Original) The method of claim 1, wherein said agent is administered along with a carrier.
- 13. (Original) The method of claim 12, wherein said carrier comprises a collagen gel, hyaluronate, alginate, calcium phosphate, polyol, or demineralized bone matrix.
 - 14. (Original) The method of claim 1, wherein said agent is administered in a matrix.
- 15. (Original) The method of claim 15, wherein said matrix comprises collagen, fibrin tissue, an endoneural sheath.
 - 16. (Original) The method of claim 15, wherein said matrix is porous.
- 17. (Original) The method of claim 1, wherein said agent is administered along with an osteogenic polypeptide.
- 18. (Original) The method of claim 17, wherein said osteogenic polypeptide is BMP-2.
 - 19. (Original) The method of claim 1, wherein said bone is metaphyseal bone.
- 20. (Original) The method of claim 19, wherein said metaphyseal bone is primal femur, proximal humerus, distal radius or vertebral body.
- (Original) A method for reducing the incidence of a bone fracture in a subject, the method comprising administering to a site at risk of bone fracture in said subject a therapeutically effective amount of an agent that inhibits BMP-3 activity.

- 22. (Original) A method for treating osteoporosis in a subject, the method comprising the method comprising administering to said subject therapeutically effective amount of an agent that inhibits BMP-3 activity in said host.
- 23. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and an agent that, when introduced into a host, results in inhibition of expression of a BMP-3 gene or activity of a BMP-3 polypeptide in said host.
- 24. (Original) The pharmaceutical composition of claim 23, wherein said agent is a nucleic acid that inhibits expression of a BMP-3 gene in said host.
- 25. (Original) The pharmaceutical composition of claim 23, wherein said agent is a BMP-3 antibody.
- 26. (Original) The pharmaceutical composition of claim 23, further comprising a carrier.
- 27. (Original) The pharmaceutical composition of claim 23, further comprising a matrix.

Claims 28-31 (Cancelled)

32. (Original) A method of antagonizing BMP-2 activity in host, the method comprising administering to said subject an agent that increases activity of BMP-3 in said host.

Claims 33-38 (Cancelled)